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FILE COVERS 1907 - 4 Dec 2008 VOL 149 ISS 23

FILE LAST UPDATED: 3 Dec 2008 (20081203/ED)

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=> s 9068-38-6 (s) (mn or manganese)

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...

Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L2 12323 L1

451777 MN

5447 MNS

454993 MN

(MN OR MNS)

413054 MANGANESE

113 MANGANESES

413065 MANGANESE

(MANGANESE OR MANGANESES)

L3 9 L2 (S) (MN OR MANGANESE)

=> s L3 and (inhibit### or modulat### or increas### or decreas### or elevat###)
2090897 INHIBIT###

411705 MODULAT###
 4604519 INCREAS###
 2568123 DECREAS###
 384482 ELEVAT###

L4 3 L3 AND (INHIBIT### OR MODULAT### OR INCREAS### OR DECREAS### OR ELEVAT###)

=> s L4 and py<2004

24012945 FY<2004

L5 2 L4 AND PY<2004

=> d L5 ibib abs 1-2

L5 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:757875 CAPLUS <<LOGINID::20081204>>

DOCUMENT NUMBER: 139:272921

TITLE: Manganese ion regulation of reverse transcriptase activity, methods of modulating reverse transcriptase, and drug screening and therapeutic use
 INVENTOR(S): Boeke, Jef D.; Bolton, Eric C.

PATENT ASSIGNEE(S): The Johns Hopkins University School of Medicine, USA

SOURCE: PCT Int. Appl., 61 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003078650	A2	20030925	WO 2003-US7879	20030312 <--
WO 2003078650	A3	20040311		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2003220277	A1	20030929	AU 2003-220277	20030312 <--
US 20050123624	A1	20050609	US 2005-507252	20050128
PRIORITY APPLN. INFO.:			US 2002-363708P	P 20020312
			WO 2003-US7879	W 20030312

AB Methods of identifying agents that modulate reverse transcriptase activity in a cell by affecting manganese ion transport across a membrane of the cell are provided, as are agents identified using such methods. High throughput screening assay for agents that alter manganese transporter activity is disclosed. Also provided are methods of modulating reverse transcriptase activity by effecting manganese ion concentration. In addition, methods of reducing or inhibiting infection of cells with a retrotransposable element are provided.

L5 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2002:351336 CAPLUS <<LOGINID::20081204>>

DOCUMENT NUMBER: 137:75669

TITLE: Inhibition of reverse transcription in vivo by elevated manganese ion concentration

AUTHOR(S): Bolton, Eric C.; Mildvan, Albert S.; Boeke, Jef D.

CORPORATE SOURCE: Department of Molecular Biology and Genetics, The
 Johns Hopkins University School of Medicine,
 Baltimore, MD, 21205, USA

SOURCE: Molecular Cell (2002), 9(4), 879-889
 CODEN: MOCEFL; ISSN: 1097-2765

PUBLISHER: Cell Press

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Mutations in PMR1, a yeast gene encoding a calcium/manganese exporter,
 dramatically decrease Ty1 retro-transposition. Ty1 cDNA is
 reduced in pmr1 mutant cells, despite normal levels of Ty1 RNA and
 proteins. The transposition defect results from Mn2+ accumulation that
 inhibits reverse transcription. Cytoplasmic accumulation of Mn2+
 in pmr1 cells may directly affect reverse transcriptase (RT) activity.
 Trace amts. of Mn2+ potentially inhibit Ty1 RT and HIV-1 RT in
 vitro when the preferred cation, Mg2+, is present. Both Mn2+ and Mg2+
 alone activate Ty1 RT cooperatively with Hill coeffs. of 2, providing
 kinetic evidence for a dual divalent cation requirement at the RT active
 site. We propose that occupancy of the B site is the major determinant of
 catalytic activity and that Mn2+ at this site greatly reduces catalytic
 activity.

REFERENCE COUNT: 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS
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